

Comparative Anthelmintic Activity of Different Extracts of *Portulaca Oleraceae* L. Whole Plant

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ABSTRACT

Parasitic diseases, especially, helmenthiasis pose a threat to human health and livestock alike. This imposes a greater burden of health care expenditure in tropical countries where these are endemic. Recently, the research for naturally- derived anthelmintics has been encouraged. *Portulaca oleracea*, commonly called as Purslane, has been part of diet in Asian and Mediterranean region. In the present study, various extracts of Purslane were tested for their anthelmintic activity on Indian adult earthworms (*Pheretima posthuma*). The anthelmintic activity of all the extracts was concentration- dependent and comparable to that of standard; while aqueous extract exhibited greatest activity. Further in- vivo tests and phytochemical research would shed light on potential activities of Purslane.


Keywords: *Portulaca oleracea*, anthelmintic, *Pheretima posthuma*.

INTRODUCTION

Infections by helminthes are a major cause of concern for human health and cattle. Larvae of helminthes predominantly exist in the intestinal tract leading to malnutrition, pneumonia, eosinophilia and anemia¹. Other clinical manifestations include

dysentery, dermatological disorders, loss of appetite and loss of body weight. Cases of presence of eggs or larvae in other tissues in the body have also been reported¹⁻². Parasitic diseases, especially, helmenthiasis are more common in the tropical countries including Asian countries and are endemic in nature³. Repeated and unregulated use



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of chemical anthelmintic drugs over considerable years has resulted in resistance amongst helminthes to these drugs. As a result, discovery of novel anthelmintic drug candidates is of prime importance to minimize the damage caused by helminthes². Owing to the general observations, naturally derived drug candidates have fewer side effects as compared to their chemically synthesized counterparts. This has lead to pharmacognostic investigations of anthelmintic activity of plant- derived products in recent years. As a primary requirement for such study, the plant candidate should be a part of accepted diet. One such potential plant candidate could be *Portulaca oleracea*, commonly called as Purslane.

Purslane is a common weed in field crops and turfgrass found in countries of Asia and Europe⁴. In parts of central Europe and Mediterranean regions it is also grown as a potherb and used in salad. As an ethnomedicinal plant, it has been used for treatment of burns, headaches, ailments related to intestine, stomach, and arthritis⁴. Purslane has been reported to contain many bioactive compounds such as flavanoids, alkaloids, coumarins, anthraquinones, catechol, saponins and tannins⁵. In the present study, we investigated the anthelmintic activity of different extracts of Purslane whole plant, in comparison to a standard.

MATERIALS AND METHODS

Plant material

Fresh whole plant (weed) of *Portulaca oleracea* was collected from our garden and authenticated by Dr. Dhanaji S. Pawar, Associate Professor, Department of Botany, M. H. Shinde Mahavidyalaya, Tisangi.

Drugs and chemicals

Mebendazole Suspension (Cipla), ethanol, methanol, butanol, carboxymethyl cellulose (CMC), all chemicals used were laboratory and analytical grade.

Preparation of plant extract

The fresh whole weed of *Portulaca oleracea* was washed with distilled water and cut into small pieces. Then 250 gram (g) weed of *Portulaca oleracea* was extracted with 800 millitre

(ml) analytical grade ethanol in soxhlet apparatus and extracted repeatedly for 48 hrs. The same procedure of extraction was followed using methanol and butanol solvents, separately. These three different extracts were dried by solvent evaporation in thermostat water bath at 50-60 °C temperature.

Another 250 g of weed of *Portulaca oleracea* was macerated with 800 ml distilled water for 48 hours (hrs). After completion of 48 hrs it was filtered to separate water extract from the marc. The extract was concentrated in thermostat water bath at 50-60 °C temperature and all the extracts were kept in desiccator for the experiment.

Selection of experimental model

Indian adult earthworms (*Pheretima posthuma*) were used during the experiment. The earthworms were collected from a local supplier. Worms were washed with normal saline to remove all fecal matter. The earthworms of 8-10 centimeter (cm) in length and 0.2 -0.5 cm width were used for all the experiment protocol. Ready availability, anatomical and physiological resemblance of (*Pheretima posthuma*) made it to be used initially for *in-vitro* evaluation of anthelmintic activity. Time for paralysis was noted either when any movement could not be observed except when the worms were shaken vigorously. Death was included when the worms lost their motility followed by white secretions and fading away of their body colour.

RESULTS AND DISCUSSION

The Indian earthworms (*Pheretima posthuma*) of nearly equal size, six in each group was taken for experiment. Four different extracts were suspended in 1% (w/v) CMC suspension prepared in normal saline water in three different concentrations.

At various concentration 25, 50,100 µg/mL comparison of time of paralysis using standard Mebendazole and various extracts showed significant difference ($F = 6.751$, $F = 18.117$, $F = 13.888$ for all F , $p < 0.001$). However, Tukey Kramer multiple comparison test, the post- hoc test revealed that time of paralysis observed using Mebendazole and aqueous extract was similar at all studied concentrations.

Table 1: Recorded time (minutes) of paralysis and death of tested earthworms (minutes) against different concentrations of standard Mebendazole and extracts ($\mu\text{g/mL}$). Time of paralysis and death expressed in mean + S.E.M.

Conc. ($\mu\text{g/mL}$)	Mebendazole		Methanolic Extract		Aqueous Extract		Ethanolic Extract		Butanolic Extract	
	Time of paralysis (minutes)	Time of death (minutes)	Time of paralysis (minutes)	Time of death (minutes)	Time of paralysis (minutes)	Time of death (minutes)	Time of paralysis (minutes)	Time of death (minutes)	Time of paralysis (minutes)	Time of death (minutes)
25	13.2 ± 0.8	18.4 ± 0.6	18.4 ± 1.7	54.4 ± 1.7	14.2 ± 0.5	46.4 ± 0.5	17.3 ± 0.5	52.8 ± 0.8	18.9 ± 0.9	62.7 ± 1.2
50	12.5 ± 0.6	16.7 ± 0.8	17.2 ± 0.4	48.2 ± 0.6	13.8 ± 0.4	44.2 ± 0.9	16.2 ± 0.4	52.2 ± 0.6	17.0 ± 0.6	60.5 ± 1.4
100	10 ± 1.3	13.7 ± 0.8	15.0 ± 0.4	42.5 ± 0.5	11.8 ± 0.4	38.4 ± 0.4	14.8 ± 0.4	50.6 ± 0.4	16.3 ± 0.5	58.9 ± 1.4

Similarly, at various concentrations 25, 50, 100 $\mu\text{g/mL}$ comparison of time of death using standard Mebendazole and various extracts showed significant difference ($F = 258.33$, $F = 333.48$, $F = 460.06$ for all F , $p < 0.001$). However, Tukey Kramer test did not show similar time taken by any studied extract in comparison to standard Mebendazole.

All the extracts exhibit concentration-dependent activity [Table 1, Fig 1]. As can be observed from the results, amongst the extracts of *Portulaca oleraceae* the aqueous extract exhibited the most anthelmintic activity with the least time required to paralyse and completely kill the earthworms at every tested concentration. Anthelmintic activity potency of the aqueous extract was followed by ethanolic, methanolic and butanolic extracts, respectively.

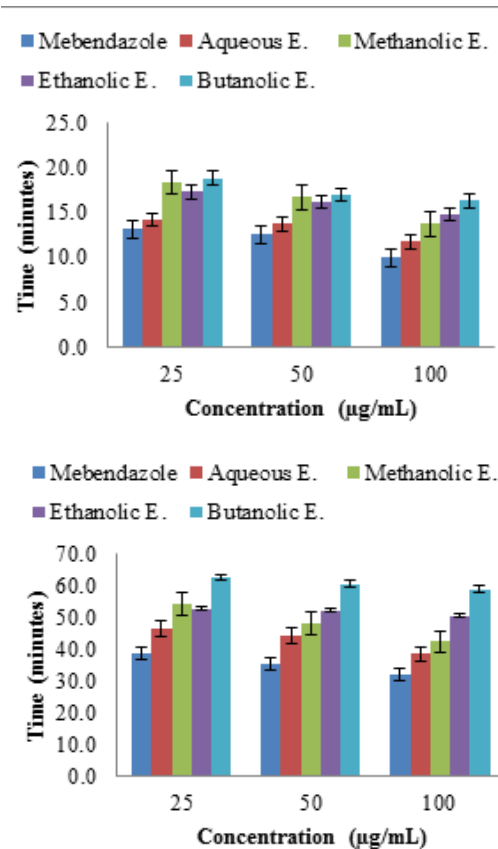


Fig. 1: Graphical representation of observed (a) time of paralysis against concentration; (b) time of death against concentration of standard Mebendazole and extracts

In a similar previously reported study, where the anthelmintic activity of different various extracts of *Euphorbia thymifolia* were tested on *Pheretima posthuma* and *Ascaris galli* worms, anthelmintic activity was observed in both aqueous and methanolic extract⁶. In another study, where anthelmintic activity of different extracts of *Enicostemma littorale* were studied, it was observed that ethanolic extract had greater paralytic activity over ethyl acetate, aqueous, chloroform, and hexane extracts⁷. The anthelmintic activity of *Luffa cylindrica* leaves extract was observed for both aqueous and methanolic extracts against *P. posthuma*². Most of the research groups have estimated that the greater amount of polyphenolics in their respective alcoholic extracts may be responsible for the observed anthelmintic activity.

CONCLUSION

After applying the Tukey Kramer multiple comparison test for the observed values, it can be concluded that the aqueous extract of *Portulaca oleraceae* exhibited anthelmintic activity similar to the standard Mebendazole with respect to time of paralysis at the tested concentrations. In order to further investigate the anthelmintic potential of this extract, phytochemical analysis and *in-vivo* tests may be undertaken.

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